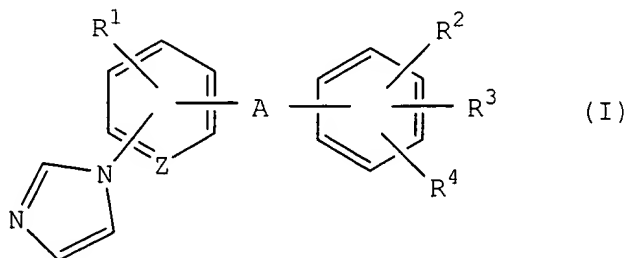


### Amendments to the Claims

1. (Currently amended) ~~A MAG-expression promoter~~ method of promoting expression of MAG comprising administering a compound of the formula (I)



wherein

$R^1$  is a hydrogen atom, a halogen atom, an alkyl group or an alkoxy group;

$R^2$  and  $R^3$  are the same or different and each is a hydrogen atom or an alkyl group;

$R^4$  is an alkyl group,  $-COOH$ ,  $-COOR^5$ ,  $-CONR^6R^7$ ,  $-CH_2NR^6R^7$ ,  $-CH_2OH$  or  $-CH_2OR^8$ ;

wherein  $R^5$  and  $R^6$  -  $R^8$  are each an alkyl group, and  $R^6$  and  $R^7$  are the same or different and each is a hydrogen atom or an alkyl group, or  $R^6$  and  $R^7$  in combination form imidazole together with the adjacent nitrogen atom;

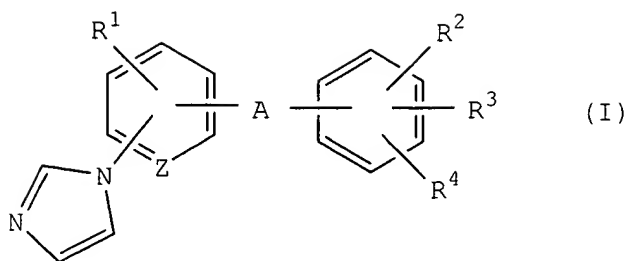
A is  $-CH(OH)-$ ,  $-C(=O)-$  or  $-CH_2-$ ; and

Z is  $=CH-$  or  $=N-$ ,

an optically active form thereof or a pharmaceutically acceptable salt thereof to a mammal.

- 2-9. (Cancel)

10. (Currently amended) ~~A method for prophylaxis and/or therapy of a disease caused by hypomyelination~~ promoting a myelination of axon, which method comprises administering a compound of the formula (I)



wherein

$R^1$  is a hydrogen atom, a halogen atom, an alkyl group or an alkoxy group;

$R^2$  and  $R^3$  are the same or different and each is a hydrogen atom or an alkyl group;

$R^4$  is an alkyl group,  $-\text{COOH}$ ,  $-\text{COOR}^5$ ,  $-\text{CONR}^6\text{R}^7$ ,  $-\text{CH}_2\text{NR}^6\text{R}^7$ ,  $-\text{CH}_2\text{OH}$  or  $-\text{CH}_2\text{OR}^8$ ;

wherein  $R^5$  and  $R^6$  are each an alkyl group, and  $R^6$  and  $R^7$  are the same or different and each is a hydrogen atom or an alkyl group, or  $R^6$  and  $R^7$  in combination form imidazole together with the adjacent nitrogen atom;

A is  $-\text{CH}(\text{OH})-$ ,  $-\text{C}(=\text{O})-$  or  $-\text{CH}_2-$ ; and

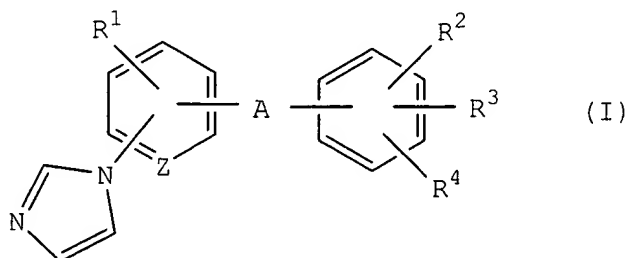
Z is  $=\text{CH}-$  or  $=\text{N}-$ ,

an optically active form thereof or a pharmaceutically acceptable salt thereof to ~~mammals~~ inclusive of human a mammal.

11. (Original) The method of claim 10, wherein, in the formula (I),  $R^1$  is a halogen atom, an alkyl group or an alkoxy group.

12. (Currently amended) A method for ~~prophylaxis and/or therapy of a disease caused by hypomyelination~~ promoting a myelination of axon, which method comprises administering 4-[ $\alpha$ -hydroxy-5-(1-imidazolyl)-2-methylbenzyl]-3,5-dimethylbenzoic acid, an optically active form thereof or a pharmaceutically acceptable salt thereof to ~~mammals~~ inclusive of human a mammal.

13. (Currently amended) A method for ~~prophylaxis and/or therapy of a disease~~ mainly presenting dysmyelination or demyelination promoting a myelination of axon, which method comprises administering a compound of the formula (I)



wherein

$R^1$  is a hydrogen atom, a halogen atom, an alkyl group or an alkoxy group;

$R^2$  and  $R^3$  are the same or different and each is a hydrogen atom or an alkyl group;

$R^4$  is an alkyl group,  $-\text{COOH}$ ,  $-\text{COOR}^5$ ,  $-\text{CONR}^6\text{R}^7$ ,  $-\text{CH}_2\text{NR}^6\text{R}^7$ ,  $-\text{CH}_2\text{OH}$  or  $-\text{CH}_2\text{OR}^8$ ;

wherein  $R^5$  and  $R^6$  are each an alkyl group, and  $R^6$  and  $R^7$  are the same or different and each is a hydrogen atom or an alkyl group, or  $R^6$  and  $R^7$  in combination form imidazole together with the adjacent nitrogen atom;

A is  $-\text{CH}(\text{OH})-$ ,  $-\text{C}(=\text{O})-$  or  $-\text{CH}_2-$ ; and

Z is  $=\text{CH}-$  or  $=\text{N}-$ ,

an optically active form thereof or a pharmaceutically acceptable salt thereof to ~~mammals inclusive of human~~ a mammal.

14. (Original) The method of claim 13, wherein, in the formula (I),  $R^1$  is a halogen atom, an alkyl group or an alkoxy group.

15. (Currently amended) A method for ~~prophylaxis and/or therapy of a disease~~ mainly presenting dysmyelination or demyelination promoting a myelination of axon, which method comprises administering 4-[ $\alpha$ -hydroxy-5-(1-imidazolyl)-2-methylbenzyl]-3,5-dimethylbenzoic acid, an optically active form thereof or a pharmaceutically acceptable salt thereof to ~~mammals inclusive of human~~ a mammal.

16-18. (Cancel)

19-30. (Cancelled)

31-36. (Cancel)

37. (New) The method of claim 10, wherein the mammal is a human.

38. (New) The method of claim 12, wherein the mammal is a human.

39. (New) The method of claim 13, wherein the mammal is a human.

40. (New) The method of claim 15, wherein the mammal is a human.